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In the Claims

Please amend the claims under the provisions of 37 C.F.R. § 1.121(c), as revised by 68-Fed. Reg. 38611 (June 30, 2003), as follows:

- 1. (Currently Amended) An isolated human *ABC1* promoter that directs transcription of a heterologous coding sequence positioned downstream therefrom, wherein the promoter is selected from the group consisting of:
 - (a) a promoter comprising nucleotides having the nucleotide sequence shown in SEQ ID NO: 1;
 - (b) a promoter comprising nucleotides having the nucleotide sequence beginning at bp -469 and ending at bp +101 624 and ending at bp 1197 of SEQ ID NO: 1; and
 - (c) a promoter comprising nucleotides having the nucleotide sequence beginning at bp $\frac{-101}{1000}$ and $\frac{-101}{1000}$ of SEQ ID NO: 1.
- (Original) The promoter of claim 1, wherein the promoter comprises the nucleotide sequence shown in SEQ ID NO: 1.
- 3. and 4. (canceled)
- 5. (Original) A recombinant expression construct effective in directing the transcription of a selected coding sequence which comprises:
 - (a) a human ABC1 promoter nucleotide sequence according to claim 1; and

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- (b) a coding sequence operably linked to the promoter, whereby the coding sequence can be transcribed and translated in a host cell, and the promoter is heterologous to the coding sequence.
- 6. (Original) The recombinant expression construct of claim 5, wherein the coding sequence encodes a transporter polypeptide.
- 7. (Original) The recombinant expression construct of claim 6, wherein the transported polypeptide is ABCA1 transmembrane transporter protein.
- 8. (Previously Presented) The recombinant expression construct of claim 6, further comprising a nucleic acid segment encoding a transactivator protein that upregulates the ABC1 promoter.
- 9. (Original) The recombinant expression construct of claim 8, wherein the transactivator protein is the Liver-X-Receptor, the Retinoid-X-Receptor, or a heterodimer of the Liver-X-Receptor and the Retinoid-X-Receptor.
- 10. (Previously Presented) A host cell in cell culture comprising the recombinant expression construct of claim 5.
- 11. (Previously Presented) The host cell of claim 10, wherein the host cell is stably transformed with the recombinant expression construct.
- 12. (Original) The host cell of claim 10, wherein the host cell is a macrophage.

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13. (Original) The host cell of claim 10, wherein the host cell is an immortalized cell.

- 14. (Original) The host cell of claim 10, wherein the cell is selected from the group consisting of RAW cells, African green monkey CV-1 cells and human 293 cells.
- 15. (Currently Amended) A method for expressing a foreign DNA in a host cell <u>in cell culture</u> comprising: introducing into the host cell <u>in cell culture</u> a gene transfer vector comprising the *ABC1* promoter according to claim 1 operably linked to the foreign DNA encoding a desired polypeptide or RNA, wherein said foreign DNA is expressed.
- 16. (Previously Presented) The method of claim 15, wherein the promoter nucleotide sequence is identical to the sequence set forth in SEQ ID NO: 1.
 - 17. (canceled)
 - 18. (Original) The method of claim 15, wherein the gene transfer vector encodes and expresses a reporter molecule.
 - 19. (Original) The method of claim 18, wherein the reporter molecule is selected from the group consisting of betagalactosidase, beta-glucuronidase, luciferase, chloramphenicol acetyltransferase, neomycin phosphotransferase, and guanine xanthine phosphoribosyltransferase.
 - 20. (Previously Presented) The method of claim 15, wherein the introducing is carried out by adenovirus infection, liposome-mediated transfer, topical application to the cell, or microinjection.

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- 21. (Original) The method of claim 15, further comprising introducing into the cell a gene transfer vector comprising a nucleic acid segment encoding a transactivator protein capable of upregulating the ABC1 promoter.
- 22. (Original) The method of claim 21, wherein the transactivator protein is the Liver-X-Receptor, the Retinoid-X-Receptor, or a heterodimer of the Liver-X-Receptor and the Retinoid-X-Receptor.
- 23. (Original) The method of claim 15, further comprising contacting the cell with a transactivator protein capable of upregulating the ABC1 promoter
- 24. (Original) The method of claim 23, wherein the transactivator protein is the Liver-X-Receptor, the Retinoid-X-Receptor, or a heterodimer of the Liver-X-Receptor and the Retinoid-X-Receptor.
- 25. (Original) The method of claim 24, further comprising contacting the cell with an agonist of the Liver-X-Receptor, of the Retinoid-X-Receptor, or of a heterodimer of the Liver-X-Receptor and the Retinoid-X-Receptor.

26-49. (canceled)

50. (Currently Amended) The isolated promoter of claim 1, wherein the promoter comprises nucleotides having the nucleotide sequence beginning at bp $\frac{101}{32}$ and ending at bp $\frac{105}{32}$ of SEQ ID NO: 1.

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51. (Previously Presented) A recombinant expression construct which comprises the nucleic acid according to claim 50.